

California Drug Recall Information



Recall Name

Hospira Recalls Magnesium Sulfate Injection Due to Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
4/13/16	50% Magnesium Sulfate Injection, USP (20mL single-dose vials) NDC: 0409-2168-02	Hospira, Inc. Lake Forest, IL	Due to the presence of a particulate within a single-dose glass fliptop vial.
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Lot: 50-343-DK Expiry: 1FEB2017	CA, nationwide	Expiry 1FEB2017

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm496111.htm